Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of the claims in the application:

Listing of Claims:

Claim 1 (currently amended): An immunogenic conjugate comprising [[a]] a synthetic homopolymer polypeptide of poly- γ -glutamic acid (γ PGA) polypeptide covalently linked to a carrier, wherein the conjugate elicits an immune response in a subject.

Claim 2 (previously presented): The conjugate of claim 1, wherein the conjugate comprises a γ PGA polypeptide comprising 5-20 glutamic acid residues.

Claim 3 (original): The conjugate of claim 1, wherein the conjugate comprises a γ PGA polypeptide comprising 10-15 glutamic acid residues.

Claim 4 (original): The conjugate of claim 1, wherein the conjugate comprises a decameric γ PGA polypeptide.

Claim 5 (currently amended): The conjugate of claim 1, wherein the carrier is selected from the group consisting of: (a) bovine serum albumin, (b) (a) recombinant B. anthracis protective antigen, (e) (b) recombinant P. aeruginosa exotoxin A, (d) (c) tetanus toxoid, (e) (d) diphtheria toxoid, (f) (e) pertussis toxoid, (g) (f) C. perfringens toxoid, (h) (g) hepatitis B surface antigen, (i) (h) hepatitis B core antigen, (j) (i) keyhole limpet hemocyanin, (k) (j) horseshoe crab hemocyanin, (l) (k) edestin, (m) (l) mammalian serum albumins, (n) (o) mammalian immunoglobulins, analogs or mimetics of (a) (n), and (p) combinations of two or more thereof.

Claim 6 (original): The conjugate of claim 1, wherein the carrier comprises recombinant *B. anthracis* protective antigen.

Claim 7 (canceled).

Claim 8 (currently amended): The conjugate of claim 1, wherein the poly-γ-glutamic acid (γPGA) polypeptide-comprises the D- or L-conformation is the D-conformation, the L-conformation, or a mixture of the D-conformation and the L-conformation.

Claim 9 (currently amended): The conjugate of claim 1, wherein the poly- γ -glutamic acid (γ PGA) polypeptide-comprises is a γ DPGA polypeptide.

Claim 10 (currently amended): The conjugate of claim 1, wherein the poly- γ -glutamic acid (γ PGA) polypeptide comprises is a decameric γ DPGA polypeptide and the carrier comprises recombinant *B. anthracis* protective antigen.

Claim 11 (previously presented): The conjugate of claim 1, wherein the carrier is covalently linked to either the amino or carboxyl terminus of the poly- γ -glutamic acid (γ PGA) polypeptide.

Claim 12 (previously presented): The conjugate of claim 1, wherein the carrier is covalently linked to the poly- γ -glutamic acid (γ PGA) polypeptide via a thioether, disulfide, or amide bond.

Claim 13 (previously presented): The conjugate of claim 1, wherein the density of poly- γ -glutamic acid (γ PGA) polypeptide to carrier is between about 5:1 and about 32:1.

Claim 14 (previously presented): The conjugate of claim 1, wherein the density of poly- γ -glutamic acid (γ PGA) polypeptide to carrier is between about 10:1 and about 15:1.

Claim 15 (original): The conjugate of claim 1, wherein the γ PGA polypeptide is covalently linked to the carrier via an aldehyde (CHO)/adipic acid hydrazide (AH) linkage.

Claim 16 (currently amended): A composition comprising the conjugate of claim 1 and a pharmaceutically acceptable carrier vehicle.

Claim 17 (original): The composition of claim 16, further comprising an adjuvant.

Claim 18 (currently amended): A composition comprising the conjugate of claim 9 and a pharmaceutically acceptable earrier vehicle.

Claim 19 (original): The composition of claim 18, further comprising an adjuvant.

Claim 20 (currently amended): A method of eliciting an immune response against a *Bacillus* antigenic epitope in a subject, comprising introducing into the subject the composition of claim 17 16, thereby eliciting an immune response in the subject.

Claim 21 (currently amended): The method of claim 20, wherein the immune response is elicited against the *Bacillus* capsular poly- γ -glutamic acid (γ PGA) polypeptide.

Claim 22 (currently amended): The method of claim 20, wherein the immune response is elicited against the *Bacillus* capsular poly- γ -glutamic acid (γ PGA) polypeptide and the carrier.

Claims 23-33 (canceled).

Claim 34 (currently amended): An immunogenic conjugate comprising a *Bacillus* eapsular poly- γ -glutamic acid (γ PGA) polypeptide covalently linked to a carrier, wherein the carrier is selected from the group consisting of: (a) recombinant *B. anthracis* protective antigen, (b) recombinant *P. aeruginosa* exotoxin A, (c) tetanus toxoid, (d) diphtheria toxoid, (e) pertussis toxoid, (f) *C. perfringens* toxoid, (g) hepatitis B surface antigen, (h) hepatitis B core antigen, (i) keyhole limpet hemocyanin, (j) horseshoe crab hemocyanin, (k) edestin, (l) mammalian serum albumins, analogs or mimetics of (a)-(l), and (m) combinations thereof, and wherein the conjugate elicits an immune response in a subject.

Claim 35 (previously presented): The conjugate of claim 34, wherein the carrier comprises recombinant *B. anthracis* protective antigen.

Claim 36 (currently amended): The conjugate of claim 34, wherein the *Bacillus* eapsular γ PGA polypeptide comprises a *B. anthracis*, *B. licheniformis*, *B. pumilus*, or *B. subtilis* γ PGA polypeptide.

Claim 37 (currently amended): The conjugate of claim 34, wherein the *Bacillus* eapsular γ PGA polypeptide comprises the D- or L-conformation is the D-conformation, the L-conformation, or a mixture of the D-conformation and the L-conformation.

Claim 38 (currently amended): The conjugate of claim 34, wherein the *Bacillus* eapsular γ PGA polypeptide eomprises is a *B. anthracis* capsular γ DPGA polypeptide.

Claim 39 (currently amended): The conjugate of claim 34, wherein the carrier is covalently linked to either the amino or carboxyl terminus of the *Bacillus* eapsular γ PGA polypeptide.

Claim 40 (currently amended): The conjugate of claim 34, wherein the carrier is covalently linked to the *Bacillus* eapsular γ PGA polypeptide via a thioether, disulfide, or amide bond.

Claim 41 (currently amended): The conjugate of claim 34, wherein the *Bacillus* capsular γPGA polypeptide is covalently linked to the carrier via an aldehyde (CHO)/adipic acid hydrazide (AH) linkage.

Claim 42 (currently amended): A composition comprising the conjugate of claim 34 and a pharmaceutically acceptable earrier vehicle.

Claim 43 (previously presented): The composition of claim 42, further comprising an adjuvant.

Claim 44 (currently amended): A method of eliciting an immune response against a *Bacillus* antigenic epitope in a subject, comprising introducing into the subject the composition of claim 43 42, thereby eliciting an immune response in the subject.

Claim 45 (previously presented): The method of claim 44, wherein the immune response is elicited against the *Bacillus* capsular poly- γ -glutamic acid (γ PGA) polypeptide.

Claim 46 (previously presented): The method of claim 44, wherein the immune response is elicited against the *Bacillus* capsular poly- γ -glutamic acid (γ PGA) polypeptide and the carrier.

Claim 47 (new): The conjugate of claim 1, wherein the carrier is a polysaccharide or a polypeptide.

Claim 48 (new): The conjugate of claim 1, wherein the carrier is a bacterial toxin or a viral protein.

Claim 49 (new): The conjugate of claim 5, wherein the carrier is selected from the group consisting of (a) recombinant *B. anthracis* protective antigen, (b) recombinant *P. aeruginosa* exotoxin A, (c) tetanus toxoid, (d) diphtheria toxoid, (e) pertussis toxoid, (f) *C. perfringens* toxoid, (g) hepatitis B surface antigen, and (h) hepatitis B core antigen.

Claim 50 (new): The conjugate of claim 9, wherein the carrier is B. anthracis protective antigen, and the conjugate elicits an immune response against γ DPGA and against B. anthracis protective antigen.

Claim 51 (new): The conjugate of claim 9, wherein the conjugate elicits IgG anti- γ DPGA antibodies in the subject.

Claim 52 (new): The conjugate of claim 51, wherein the conjugate also elicits IgG anticarrier antibodies.

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Claim 53 (new): The method of claim 20, wherein the immune response elicits IgG anti-B. anthracis γ PGA antibodies and IgG anti-carrier antibodies in the subject.

Claim 54 (new): The conjugate of claim 34, wherein the carrier is selected from the group consisting of (a) recombinant *B. anthracis* protective antigen, (b) recombinant *P. aeruginosa* exotoxin A, (c) tetanus toxoid, (d) diphtheria toxoid, (e) pertussis toxoid, (f) *C. perfringens* toxoid, (g) hepatitis B surface antigen, and (h) hepatitis B core antigen.

Claim 54 (new): The conjugate of claim 38, wherein the conjugate elicits IgG anti- γ DPGA antibodies and IgG anti-carrier antibodies in the subject.

Claim 55 (new): The method of claim 44, wherein the immune response elicits IgG anti-B. anthracis γ PGA antibodies and IgG anti-carrier antibodies in the subject.

Claim 56 (new): The conjugate of claim 1, wherein the conjugate includes a plurality of γ PGA polypeptide chains per carrier molecule.

Claim 57 (new): The conjugate of claim 1, wherein the conjugate has a density of γ PGA chains to carrier molecule of at least about 5:1.

Claim 58 (new): The conjugate of claim 34, wherein the conjugate includes a plurality of γ PGA polypeptide chains per carrier molecule.

Claim 59 (new): The conjugate of claim 34, wherein the conjugate has a density of γ PGA chains to carrier molecule of at least about 5:1.

Claim 60 (new): Am immunogenic conjugate comprising poly- γ -glutamic acid (γ PGA) covalently linked to a carrier, wherein the conjugate elicits an immune response in a subject, and the conjugate has a density of γ PGA chains to carrier molecule of at least about 5:1.

Claim 61 (new): The conjugate of claim 56, wherein the carrier is a polymeric carrier.

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Claim 62 (new): The conjugate of claim 57, wherein the carrier is a polymeric carrier.